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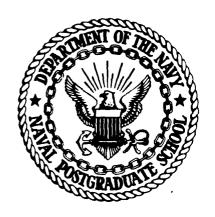
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# NAVAL POSTGRADUATE SCHOOL Monterey, California



MIXED LAYER MODELING OF AEROSOLS IN THE MARINE BOUNDARY LAYER

bу

K. L. Davidson, C. W. Fairall, and G. E. Schacher

May 1982

Technical Report

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Southern California during the CEWCOM-78 experiment. The test was with two different methods of obtaining the relevant meteorological and aerosol parameters: 1) actual measurements and 2) dynamic boundary layer model prediction and parameterization. In the first case the model reproduced the aerosol data within a factor of 1.5 while in the second case the uncertainty factor was 2.0. In either case the model only modestly outperforms the much simpler Wells-Munn-Katz (WMK) model, which uses only local specification of the wind speed and humidity. Suggested improvements of the mixed layer model are presented.

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#### 1. Introduction

This is one of a series of reports that describes the Naval Postgraduate School's (NPS) approach to marine atmospheric boundry layer (MARL) modeling. The basics of the approach and the status of modeling and parameterization of the pertinent physical processes are given in Fairall et al. (1981). The utility of the model for tactical use and initial validation of the model are given in Davidson et al. (1982). In this report we describe the use of the model for analysis and prediction of the aerosol content of the MARL.

Interest in marine aerosols has increased recently because of their contribution to the scattering and absorption of light. Estimating these influences on electro-optical (EO) system performance has been emphasized in several studies such as Barnhardt and Streete (1970). An aerosol size of interest is that associated with locally generated sea salt because of its effects on IR as well as visible wavelengths. Measured distributions in the sea salt size range show several orders of magnitude variability, and considerable effort has been expended to normalize these measurements by correcting appropriately for relative humidity and wind speed. These two meteorological quantities are considered because of their role in generation (wind), transport (wind), and growth (relative humidity) of aerosols.

Both empirical and theoretical bases exist for formulating expressions for equilibrium aerosol distributions. Past quantifications of the dependence of equilibrium distributions on relative humidity were made by Fitzgerald (1975). Dependence on turbulent transport was considered by Toba (1965), and dependence on surface generation by Chaen (1973). However, aerosol models in use today utilize only parameterizations of the effects

of relative humidity and wind speed on the equilibrium aerosol distributions (e.g., Wells et al, 1977). Recent evaluations have shown that these models are limited to mean (that is, the average aerosol density encountered at a given wind speed and humidity) distributions (Fairall et al, 1982). These models are inherently limited because some processes in the atmospheric mixed layer which affect aerosol concentrations are not considered.

In this study we will present an approach for including meteorological descriptions which encompass processes and structures of the whole marine atmospheric boundary layer. The top of the boundary layer is capped by the marine inversion where entrainment of overlying air takes place. Because entrainment mixes clear (non-marine) air into the marine layer this process is as important as surface layer aerosol fluxes in determining the equilibrium concentration. The primary goal of our examination will be to incorporate in aerosol descriptions recently established features of the inversion capped marine boundary layer. The features will be presented in terms of an integrated slab model. Such a model is especially desirable since the only input parameters needed are routine meteorological observations.

### 2. Description and Evaluation of Existing Aerosol Models

A representative example of current models for estimating equilibrium aerosol distributions is that formulated by Wells et al (1977). Their formulation was modified by investigators in the Navy Electro-Optical/Meteorology Program (Hughes, 1980). The number density spectrum, n(r), of the sea salt component is described in the modified version as

$$n(r) = (r/a) \cdot 1.62 \cdot (C_1 + C_2 v^{\delta}) / Fe^{-Z/h_0 F - 8.5(r/a)^{\gamma}}$$
, (1)

where

r = the particle radius in  $\mu m$ 

u = the wind speed

v = 0.5 for  $u \le 4$  m/s,

v = u - 3.5 for u > 4 m/s,

 $F = 1 + (v/60)^3$ ,

 $\gamma = 0.384 - 0.00293 \text{ v}^{1.25}$ ,

Z = height above sea surface, m ,

 $h_{\rm O} = {\rm scale\ height}$ , m (800 m for Z < 1000 m) ,

 $a = 0.81 \exp (0.066S/(1.058 - S))$ ,

S = H/100, (H is relative humidity in per cent).

The other constants are

v(m/s)	<sub>c1</sub>	<sup>C</sup> 2	δ
v <u>&lt;</u> 7	350	1000	1.15
v > 7	0	<b>69</b> 00	0.29

The relative humidity growth factor, a, has the form suggested by Fitz-gerald (1975). The height dependence is exponential, with the scaling height,  $h_{\rm O}$ , being a function of wind speed as suggested by Toba (1961). The leading term  $(C_1+C_2, v^{\delta})$  corresponds to local generation and has a property of the second se

law dependence on wind speed. This particular version is referred to as the WMK model.

Examination of a considerable amount of data obtained in the Northeast Atlantic and the Eastern Pacific indicates that Eqn 1 is incomplete. Figure 1 shows the height dependence of total aerosol volume from a sample set of Eastern Pacific data. The three aerosol profiles correspond to (1) the observed sea salt aerosol volume, V, (2) the observed sea salt aerosol volume adjusted to 80% relative humidity equilibrium sizes,  $V_{\text{O}}$ , and (3) the WMK predicted volume adjusted to 80% (open circles). These data were obtained under forced convective conditions and with active local production. In order to examine only the height dependence issue, observed and model values have been matched at the surface to remove the production influence. It is clear from the figure that within the mixed layer the observed decrease of aerosol volume with height is less than that predicted by Eqn 1. The surface generated aerosols appear to be well-mixed below the inversion when normalized to remove the influence of relative humidity. This has also been observed by other investigators (Blanchard and Woodcock, 1980; Hughes, 1980; Johnson and Hering, 1981). Aerosol volume corrected for relative humidity can be considered to be reasonably well mixed from the inversion down to approximately the 10-meter level for these conditions.

Evidence supporting our assertion that existing scaling of production effects are reasonable only for mean or climatological purposes appears in results obtained in the Northeast Atlantic. This point is illustrated in Fig. 2 where normalized measured aerosol volume densities at  $r=5~\mu m$  arithmeter corresponding prediction from Eqn 1 are compared. The values and trends in the predicted and mean results are in reasonable agreement. However, the standard deviation is so large that, at a given time, only 67% of the

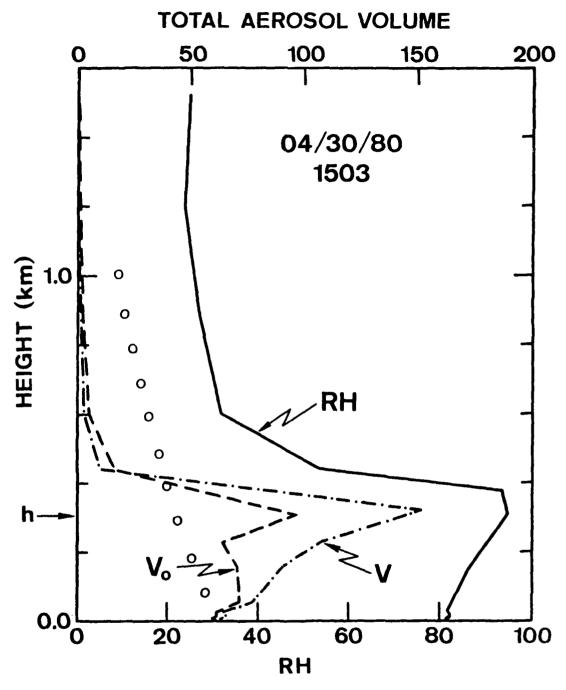


Fig. 1 Height dependence of aerosol volume and relative humidity above the ocean. The curve labeled V is the ambient aerosomototal volume  $(\mu m^3/cm^3)$ , the curve label  $V_O$  is the volume corrected to 80% reference humidity. The open circles represent the equivalent  $V_O$  from the WMK model (Eq. 1)

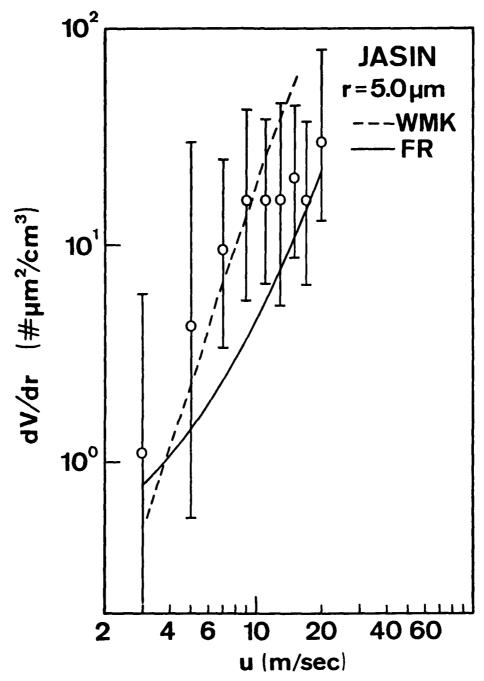


Fig. 2 Aerosol volume spectrum, dV/dr = V(r), at r = 5 µm as a function of wind speed, u. The circles represent average values obtained from measurements in the North Atlantic with the standard deviation shown as brackets. The dashed line is the WMK model and the solid line is a similar model taken from earlier measurements in the Atlantic

observed aerosol distributions will be within a factor of three of the average. The point is that no matter how accurately the model predicts the average aerosol density at a given wind speed and relative humidity, the factor of three for the RMS variation cannot be eliminated without considering more parameters than instantaneous wind speed and relative humidity.

#### Boundary Layer Model

The prime feature characterizing the marine atmospheric boundary layer is that it is convectively mixed, and hence homogeneous, up to a height h, where it is capped by an inversion. The mixed layer is cool and moist relative to the overlying stable layer (shown in Fig. 3).

The fundamental assumption of existing models of the boundary layer is that well-mixed properties remain well mixed when undergoing evolutions in time. If the value of any well-mixed property, X, were to change solely because of turbulent fluxes at the top and bottom boundaries and uniform horizontal advection, and if X remains well mixed after the change, then it follows that the vertical flux profile, <W'X'>, must be linear. For this case the appropriate balance expression for the local change of X is (Lilly, 1968)

$$dX/dt + V_{H} \cdot VX = (\langle W'X' \rangle_{O} - \langle W'X' \rangle_{i})/h , \qquad (2)$$

where  $V_{\text{H}} \cdot VX$  is the horizontal advection, the brackets represent a suitable average for the responsible turbulent scales, and subscripts o and i refer to surface and inversion values, respectively.

Difficulties in making measurements of the fluxes at the inversion and evaluating the horizontal advection make Eqn 2 impractical for single location observations without further simplification and scaling. The horizontal advection term is excluded in the following discussion not because it is necessarily negligible but simply to limit the scope of this paper. The surface flux is easily calculated using the bulk aerodynamic method. This leaves the flux at the inversion to be parameterized in order for Eqn 2 to be a usable balance expression for single station assessments.

The relationship between the equivalent flux property at the inversion  $\langle W'X' \rangle_i$ , the rate of change of the mixed layer depth (dh/dt), the mean

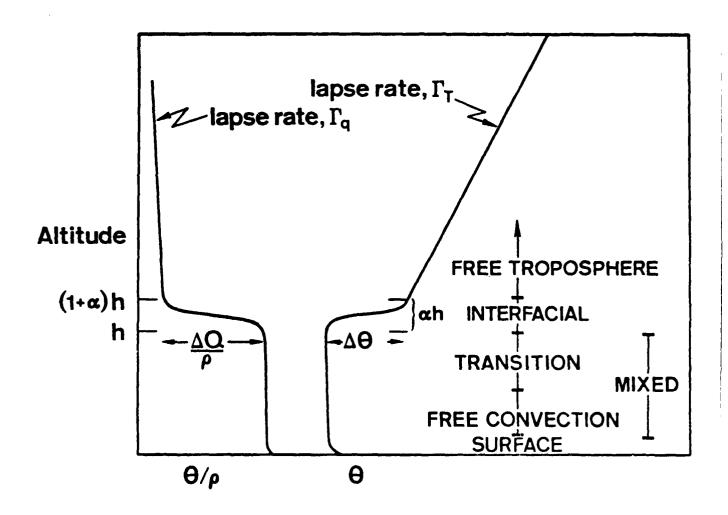


Fig. 3 Marine boundary layer well-mixed structure in a two layer idealization. The height of the mixed layer is h.  $\theta_V$  is the virtual potential temperature, Q the water vapor density and  $\rho$  the density of air

vertical motion, W, the entrainment velocity,  $W_e$ , and the "jump" in X,  $\Delta X$ , at the inversion is:

$$dh/dt = W + W_{\rho} , \qquad (3a)$$

$$\langle W'X' \rangle_{\dot{1}} = -W_e \Delta X$$
 (3b)

The utility of these expressions depends on the capability to estimate the quantity  $W_e$  from bulk parameters. An estimate of the  $\langle W'X' \rangle_i$  term in Eqn 3 can be made if changes of the inversion height and subsidence rates are known ( $W_e = dh/dt - W$ ). As such, it would not be useful for prediction since one uses  $W_e$  to predict dh/dt, but it could be applied to analytically explain contributions to observed changes in X.

All integrated schemes relate the availability and consumption of turbulent kinetic energy at the inversion to surface, cloud regime, and inversion parameters. Turbulent kinetic energy available for entrainment is derived from surface layer and cloud region buoyancy fluxes and mechanical turbulence associated with surface layer and inversion layer wind shears. The effectiveness of the available energy in driving entrainment depends on the static stability of the inversion layer.

The simplest formulation for  $W_e$  is for the cloud free zero-order model (Fig. 3) where the static stability is scaled by the jump in virtual potential temperature. The available kinetic energy is assumed to be a fraction of the <u>surface</u> buoyancy flux. The expression for  $W_e$  is (Lilly, 1968)

$$W_{e} = f \langle W' \theta_{V}' \rangle / \Delta \theta_{V} , \qquad (4)$$

where suggested values of f have ranged from .1 to .3.

The complexity of parameterizations of this type increases when cloud region bubyancy is included and when more realistic structures are used for the inversion zone. Deardorff (1978) and Stage and Businger (1981) are suggested for more complete descriptions of the scaling procedures. In all

cases, however, a useful scaling velocity for the entrainment rate is the convective mixing velocity,

$$W_{\star} = (9Q_{O}Z_{i}/T)^{1/3} , \qquad (5)$$

where  $Q_O = \langle W'\theta'_{VO}, Z_i = (1+\alpha)h$  is the inversion height, T the absolute temperature, and g the acceleration due to gravity. The equivalent expression to Eq. 4 has the form

$$W_e/W_{\star} = fW_{\star}^2/(gZ_{i}\Delta\theta_{v}/T) . \qquad (6)$$

# 4. Aerosol Considerations in a Well-Mixed Model

The first step in applying the mixed layer model to aerosol descriptions is the establishment of the well-mixed parameters. In the case of temperature (clear sky) it is the virtual potential temperature; in the case of humidity it is the water vapor mixing ratio. Although a given aerosol particle may vary in size due to changes in ambient relative humidity, the salt particle spectrum is conserved (because of the lower number concentrations of sea salt aerosols, coalescence can generally be neglected). Thus, the well-mixed property for aerosols is the dry size spectrum mixing ratio, or equivalently, the aerosol spectrum mixing ratio at some reference relative humidity. We will consider the aerosol volume spectrum

$$V(r) = 4/3\pi r^{3}n(r) , \qquad (7)$$

where n(r) = dN/dr is the number density spectrum.

We shall define V(r) as the volume of aerosol particles per cm<sup>3</sup> per radius increment at STP and at the reference saturation ratio,  $S_{\rm O}=0.8$ . At some height in the mixed layer, where the local ambient air density is  $\rho$  and the saturation is S, the aerosol volume spectrum is

$$V'(r_s(s)) = V(r_o) g^2(s) \rho/\rho_o$$
, (8)

where  $\rho_{O}$  is the density of air at the surface,  $r_{O}$  and  $r_{S}$  the particle radii at the reference and ambient saturation

$$r_{s} = r_{o}g(s) , \qquad (9)$$

and g(S) is the humidity growth factor

$$g(S) = .81 \exp(0.066 S/(1.058-S))$$
 (10)

Note that  $\rho/\rho_0$  is included to maintain a constant mixing ratio.

The next step in applying the mixed layer model to aerosols is to recognize that all aerosols do not originate locally. Both above and believe

the inversion, aerosols are advected into the local region. In terms of aerosol density, entrainment acts as an aerosol flux (out of the boundary layer) because the aerosol concentrations above and below the inversion are different. In the mixed layer model the entrainment acts on the jump across the inversion and the flux of a scalar property is

$$\langle W'X' \rangle_{i} = -W_{e} \Delta X = -W_{e} (X_{p} - X_{m}) ,$$
 (11)

where  $\mathbf{X}_{\mathbf{p}}$  is the quantity just above the inversion and  $\mathbf{X}_{\mathbf{m}}$  is the quantity just below the inversion.

With aerosols we find it useful to consider two components as distinguishable from each other. The total aerosol volume is the sum of the surface generated salt component,  $V_{\rm S}$ , and the background (continental) component,  $V_{\rm C}$  (Figure 4); then

$$V_{m} = V_{sm} + V_{cm} , \qquad Z < h \qquad (12a)$$

$$V_{p} = V_{CP}, \qquad Z > h \qquad (12b)$$

where we have assumed  $V_{sp} = 0$ .

At this point Eqn 3 can be applied to each component

$$h(dV_{sm}/dt) = \langle W'V_{s}' \rangle_{O} - (W_{e}+W_{km})V_{sm},$$
 (13a)

$$h(dV_{cm}/dt) = W_e(V_{cp}-V_{cm}) - W_{km} V_{cm} + W_{kp} V_{cp}$$
, (13b)

where we have neglected horizontal advection, assumed negligible local production for the continental component ( $\langle W'V_C' \rangle_O \approx 0$ ) and included the Stokes gravitational fallout term  $W_K$  (Wu, 1979). The fallout rates above and below the inversion are different because of the change in the aerosc spectra caused by the humidity growth factor. The Stokes velocity is calculated from

$$W_{k} = 2g(\rho_{w} - \rho)r_{O}^{2}g^{2}(S)/(9E\rho)$$
, (14)

where  $\rho_{\mathbf{w}}$  is the density of the droplet, g the gravitational acceleration.

and E the kinematic viscosity of air. Since the rate of change of the

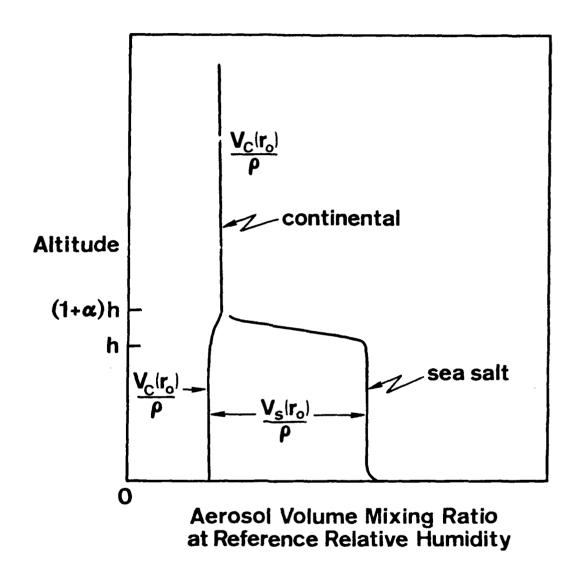


Fig. 4 Height dependence of continental,  $V_{\rm C}$ , and sea-salt,  $V_{\rm S}$ , aerosols in the mixed layer model. As shown, the aerosol volumes are represented at fixed humidity

continental component ( $dV_{\rm cm}/dt$ ) due to mixed-layer processes is often small compared to the advective effects, no attempt will be made to investigate Eqn 13b. The actual process of determining the separate components of a measured aerosol spectrum presently relies on the fact that virtually all particles with radii less than 0.1  $\mu m$  are of "continental" origin (Fairall et al, 1982).

# 5. Marine Boundary Layer Experiment

The previously described aerosol balance expression will be evaluated with aerosol and atmospheric mixed layer measurements made off the U.S. West Coast. They were made during the Cooperative Experiment on West Coast Oceanography and Meteorology (CEWCCM-78) which was conducted from 25 April to 23 May 1978 by several U.S. Navy sponsored groups. Measurements were made from the research vessel ACANIA and several shoreline radiosonde stations. The general area of the experiment is shown in Figure 5. Descriptions of synoptic scale conditions influencing the Los Angeles basin throughout the entire CEWCCM-78 period have been presented by Rosenthal et al (1979). For this evaluation we chose a period, 20-21 May, toward the end of the experiment.

The 20-21 May period was one of increasingly maritime conditions in the Los Angeles basin due to onshore flow caused by the development of a thermal trough over Southern California. This thermal trough developed in conjunction with the intensification of a Pacific high located west of Washington state.

Mixed layer changes during this period were controlled by the following factors: 1) widespread subsidence was occurring over a uniform mixed layer, 2) prevailing onshore flow reduced the effect of local land/sea circulations, 3) overcast stratus-stratocumulus contributed to the entrainment at the top of the layer, and 4) advection was not a significant factor in local changes of mixed layer depths.

Deepening of the marine, or mixed, layer during the period is evident in an acoustic sounder record which was obtained on the R/V ACANIA and incomposite profiles constructed from radiosondes launched at shoreline and ship locations (Fig. 6). The acoustic sounder record and the composite

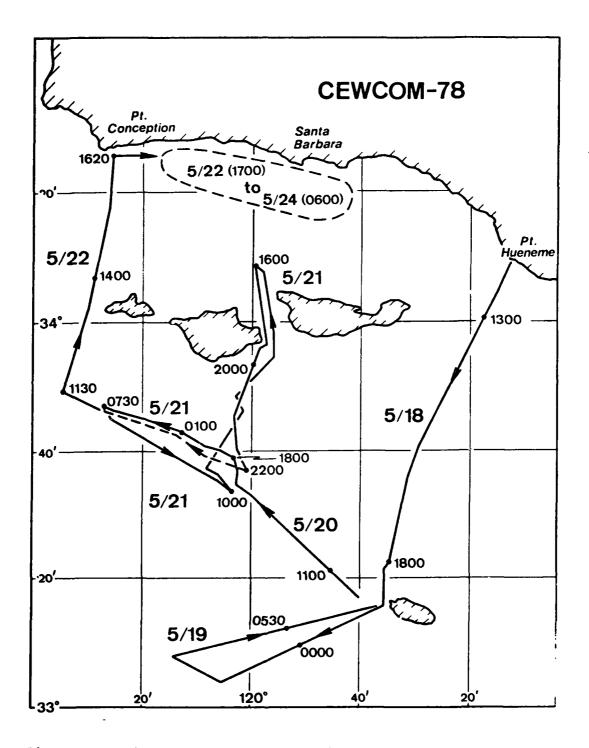


Fig. 5 Location of the CEWCOM-78 experiment in coastal southern California. The line indicates the track of the R/V ACANIA

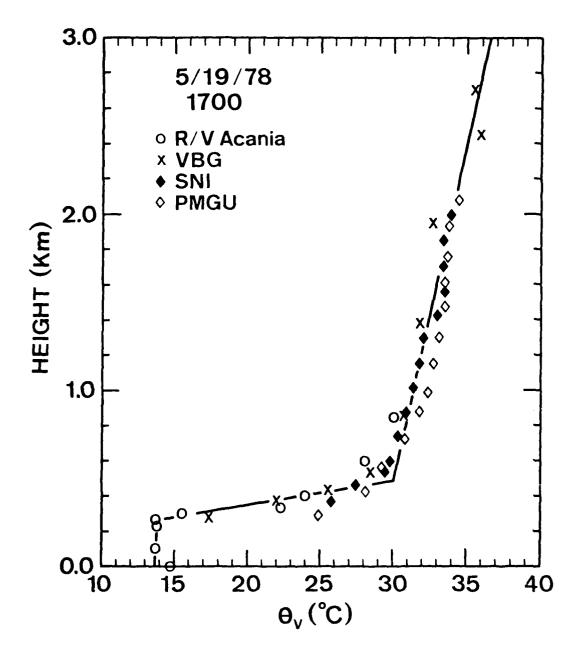


Fig. 6a Composite radiosonde measurements of  $\boldsymbol{\theta}_{_{\boldsymbol{V}}}$  at 5/19, 1700 local time

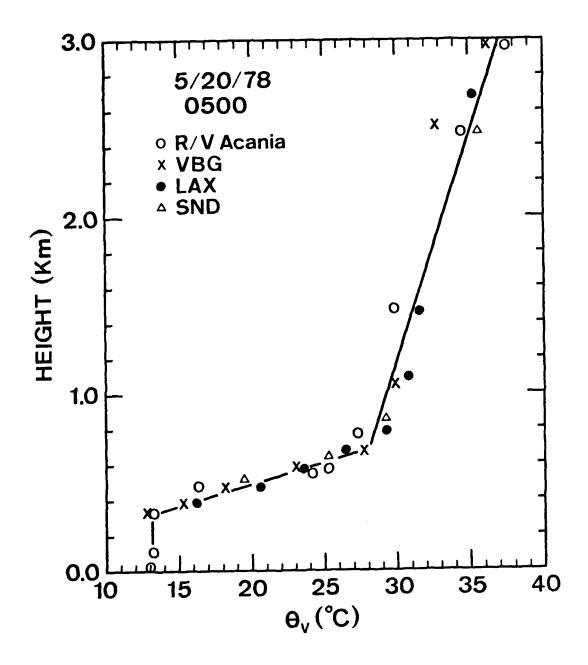


Fig. 6b Composite radiosonde measurements of  $\boldsymbol{\theta}_{_{\boldsymbol{V}}}$  at 5/20, 0500 local time

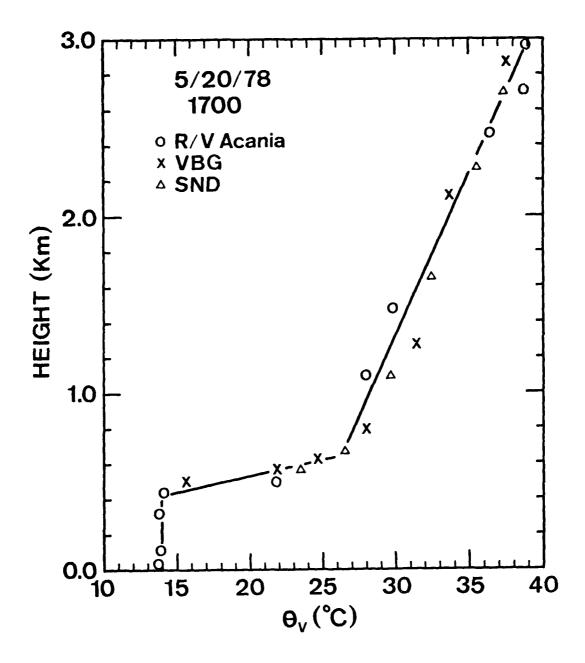


Fig. 6c Composite radiosonde measurements of  $\theta_{_{\mathbf{V}}}$  at 5/20, 1700 local time

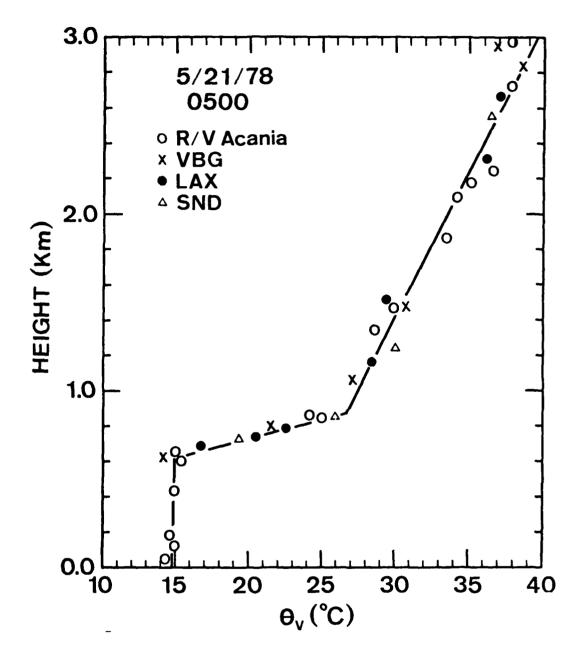


Fig. 6d Composite radiosonde measurements of  $\boldsymbol{\theta}_{V}$  at 5/21, 0500 local time .

files indicate that the mixed layer depth was nearly the same at all stations and changed uniformly.

Measurements from the R/V ACANIA provided information on the surface layer wind, temperature, humidity and aerosol spectrum. These data are summarized in Table I for the period of interest.

Table I

Meteorological surface layer data for the CEWCOM-78 analysis period.  $T_{\rm S}$  is the sea-surface temperature and T the air temperature. These are four-hour averages of half hourly observations.

Date	Time	u(m/s)	S 	T <sub>S</sub> (C)	T(C)	h( m) 
 5/20	1300	7.9	.86	14.0	12,3	425
5/20		7.9 8.7	.82	14.0	12.3	435
5/21	2100	9.8	.87	13.7	12.1	385
5/21	0100	9.2	•88	13.1	12.0	448
5/21	0500	8.2	.83	13.0	12.0	615

Practical application of a mixed layer model requires parameterization of the entrainment rate, W<sub>e</sub>. However, in Section 4 we wish to test only the aerosol part of the model, not the accuracy of various entrainment predictions. Therefore, the entrainment rate was directly estimated from the measured evolution of the well-mixed humidity and temperature over the period using the NPS boundary layer model. An entrainment rate was selected that reproduced the evolution of the non-aerosol mean variables (humidity and temperature) and then that entrainment rate was applied to the aerosol evolution.

The vertical motion, W, was calculated using the known changes in h and the values to  $W_e$  in Eqn 3a. The divergence, D, was estimated as D = -W/h (Table II).

Period	W <sub>e</sub> (aπ/sec)	Divergence (sec <sup>-1</sup> )
5/20-0500 to 5/20-1700	0.32	7.5 x 10 <sup>-6</sup>
5/20-1700 to 5/20-2400	0.32	1.3 x 10 <sup>-6</sup>
5/21-0000 to 5/21-0500	0.38	1.3 x 10 <sup>-6</sup>

#### 6. Aerosol Model Test

This section is an examination of the ability of Eqn 13a to predict evolutions of sea salt aerosol spectra given ideal estimates of the meteorological parameters. Consequently, measured (as opposed to model parameterized or predicted) values are used for non-aerosol parameters wherever possible. In particular, the wind speed, relative humidity, entrainment rate and boundary layer height for the analysis will be based on actual observations, not model predictions.

#### a. Aerosol Surface Flux

The evolution of the sea salt component of the aerosols is described by Eqn 13a where h,  $W_{\rm km}$  and  $W_{\rm e}$  are given. The only other unknown in the equation is the surface aerosol flux spectrum  $\langle W'V_{\rm S}'\rangle_{\rm O}$ . This quantity has been determined previously from data collected in the Northeast Atlantic during the JASIN experiment, (Fairall et al, 1982) and is given in Table III as a function of wind speed.

Table III Sea salt aerosol surface volume flux in  $\mu m^2/cm^2/s$ , as a function of wind speed (m/s) and particle radius ( $\mu m$ ) at S = 0.8.

U	r = 0.8	2	5	10	15	
6	1.3	1.1	2.5	1.0	0.3	_
9	4.5	3.1	4.2	3.3	2.3	
11	8.2	7.7	11.0	21.0	27.0	
13	9.1	9.2	17.0	49.0	48.0	
15	11.0	10.0	19.0	72.0	140.0	
18	17.0	11.0	24.0	92.0	180.0	

#### b. Initialization

Eqn 13a is a rate equation that relates  $dV_{sm}/dt$  to  $V_{sm}$ . Consequently the model is used to predict the evolution of  $V_{sm}$  subsequent to some initial time for which values of the relevant parameters are known or

assumed. In this example, we will use the values measured at the beginning of the analysis period. If measured aerosol values are not available, the model can be initialized with the equilibrium spectrum obtained from Eqn 13a with  $dV_{cm}/dt = 0$ :

$$V_{sm}$$
 (Equilibrium) =  $\langle W'V_{s'} \rangle_{o}/(W_{e} + W_{km})$ . (15)

# c. Results

The model was initialized for 1300 PDT on 5/20/78 with the measured spectrum. The evolution of the size spectrum was calculated with half-hour time steps at five particle sizes (r = 0.8, 2, 5, 10 and 15 µm) using the data given in Tables I, II and III. The surface flux used at each wind speed was a log interpolation of the values in Table 3 weighted by the velocity category. The results of the model calculation for three radii are shown in Figure 7 and compared with the measured data and the WMK model at four-hour intervals. Each of the three results plotted is normalized by its own particular value at the start of the analysis period. This was done to avoid biasing the comparison in favor of the mixed layer model which, since it is initialized with the measured data, would tend to automatically predict the correct results more accurately (especially in the earlier periods). The initial and final measured spectra are shown in Figure 8 along with the mixed layer and WMK model predictions.

#### d. Discussion

The mixed layer aerosol parameterization did a credible job of policiting the aerosol spectum 16 hours after initialization. A log-averaging yielded an average of the predicted to the measured spectra of 0.92 x  $^{\circ}$  1.54 (1.54 =  $e^{\sigma}$  where  $\sigma = 0.43$  is the standard deviation of the log  $\sigma$  ratio and  $\langle x \rangle x$  or  $e^{\sigma} = \langle x \rangle e^{\pm \sigma} = e^{\langle x \rangle \pm \sigma}$ ). In this case, the WW also performed well with a log-average ratio of prediction to measure

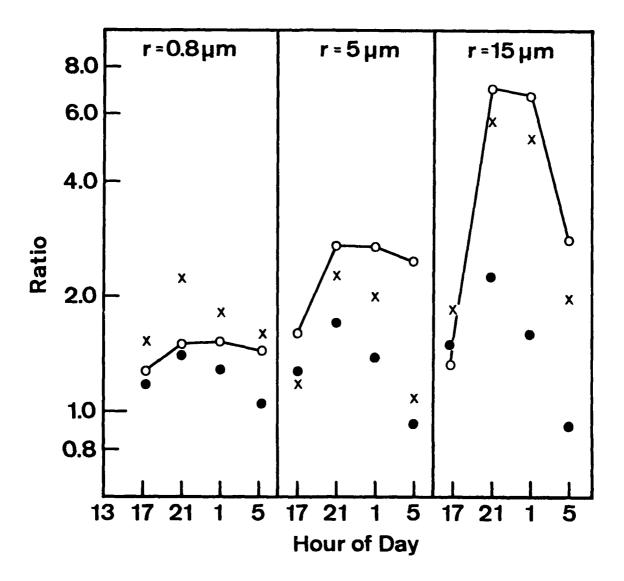


Fig. 7 Ratio of ambient aerosol volume at the hour indicated to the volume at the beginning of the period,  $V_{sm}(r,t)/V_{sm}(r,o)$ . The open circles are the mixed layer dynamic model, the X's are the measurements and the solid circles are the WMK model. The mixed layer model and measurements were normalized by the initial value of the data. The WMK ratio was obtained by normalizing with the WMK model value obtained from the initial wind speed and relative humidity

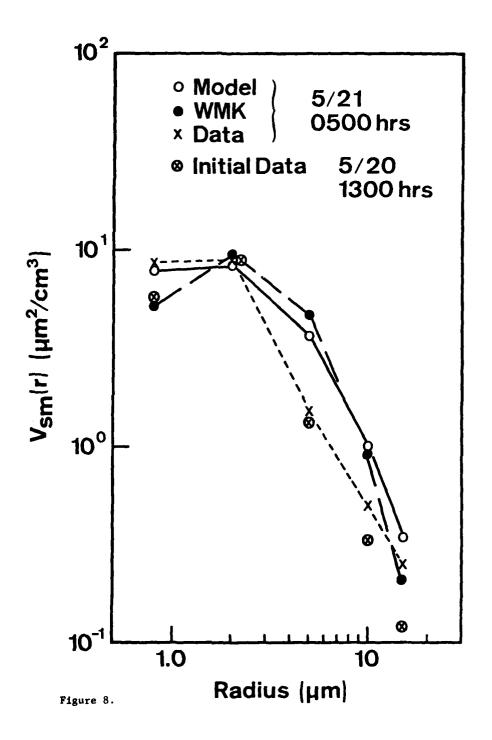


Fig. 8 Initial (circled X's) and final (X's) measured aerosc spectra. The mixed layer model spectrum is the open circ and the WMK model spectrum is the solid circles

ratios of 1.69 x or  $\div$  1.6. This is uncharacteristically good performance for the WMK type of model. In practical application, the mixed layer model will probably not perform this well since several important model parameters ( $W_e$ , h, u and S) were not model predicted but obtained from the measurements. This issue will be examined in section 8. Other sources of error (such as advection) will also have an impact. The most important parameter, the surface flux of aerosols (Table III), was based primarily on a single data set obtained in the North Atlantic. Further refinements in the surface flux can, at least potentially, greatly improve the model accuracy. The surface flux data also need to be extended to larger particle sizes.

# 7. Full Dynamic Model Test

Eqn 13a of Section 4 was tested for a 20-hour time period using measured or directly inferred values of the relevant model parameters. The aerosol model spectrum was initialized at the beginning of the analysis period with the measured spectrum. This section describes a simulation where a complete dynamical, cloud-topped boundary layer model of the Deardorff (1976) type is used to provide the necessary parameters for Eqn 13a.

#### a. Procedure

The full mixed layer model was run in three 12-hour blocks beginning at 1700 local time on 5/19/78 with subsequent re-initialization of all meteorological variables at the 0000 and 1200 GMT radiosondes (1700 and 0500 local time). The aerosol spectrum was initialized at 1700 on 5/19 using Eqn 15 and was not subsequently reset. The evolution of the aerosol spectrum was calculated using Eqn 13a with all parameters taken from the dynamic model output. The measured aerosol data were available for the last 20 hours of the 38-hour model simulation (the final time period was run for 14 hours to cover the last two hours of aerosol data). Other details of relevance are:

- The surface fluxes were obtained using standard bulk aerodynamic parameterizations.
- 2) The sea-surface temperature was assigned as the measured value at the beginning of a 12-hour forecast period and assumed to 1 constant throughout that period.
- 3) The measured wind speed for a 12-hour period was simplified two or three continuous linear segments in an effort to 3' late an accurate wind speed forecast.

- 4) The subsidence rate for the start of the test was based on an assumed climatological divergence of  $D = 1 \times 10^{-5} \text{ sec}^{-1}$ . The difference between the predicted and measured values of h at
- the end of each 12-hour period was used to re-estimate the appropriate subsidence rate for the period. This new subsidence value was then applied to the next 12-hour period.
  - 5) Advection of temperature, water vapor, mixed-layer depth and aerosols were assumed to be negligible.

#### b. Aerosol Results

Mixed layer model and WMK model aerosol spectra were calculated as described in Sections 5.1 and 4.3. The results are shown in Fig. 9 for the entire 38 hour period. Note that in this case the actual spectral values are plotted rather than the model/initial ratios shown in Fig. 7. The mixed layer model was not a significant improvement over the WMK for this data set. A log-averaging yielded an average ratio of mixed layer model to measured aerosol spectra of 0.76 x or  $\div$  2.0 while for the WMK model the ratio was 0.87 x or  $\div$  2.2. In other words, the mixed layer model predictions were typically within a factor of 2.0 while the WMK predictions were typically within a factor of 2.2. These can be contrasted with the results of Section 4.4 where, using measured parameters and spectral initialization, the mixed layer model was typically within a factor of 1.5 and the WMK was within a factor of 1.6. Thus, a more realistic application of the model doubled the uncertainty (note that a factor of 1.0 is no uncertainty).

## c. Mixed Layer Dynamics

The practical application of the full dynamic model (Section 5) led to an increase of uncertainty in the aerosol prediction from a factor of

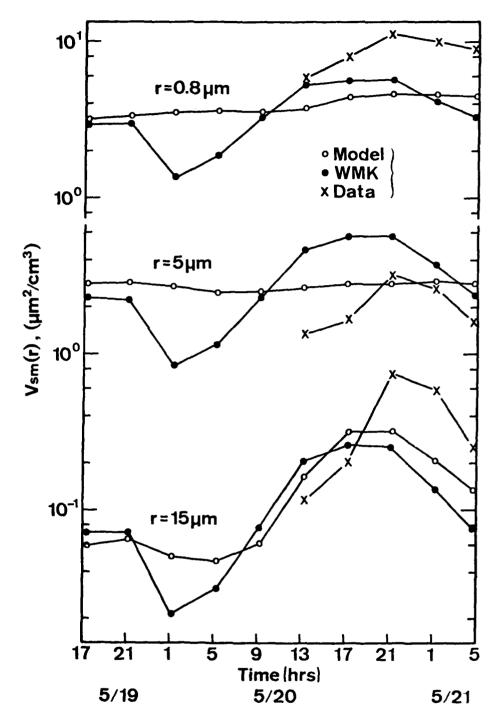


Fig. 9 Temporal evolution of the aerosol volume at  $r=0.8~\mu m$ , 5  $\mu m$  and 15  $\mu m$  over the 38 hour period beginning at 1700 local time on 5/19. The open circles are the full mixed layer model, the solid circles are the WMK model and the X's are the data

1.5 to a factor of 2. The primary sources of the increase in error are:

- 1) Use of Eqn 15 for initialization.
- 2) Model calculations of W<sub>a</sub>.
- 3) Linear smoothing of the wind speed temporal evolution.
- 4) Model calculation of h.

A comparison of model and measured values of  $W_e$ , h and W is shown in Fig. 10. Recall that the model was initialized at 12 hour intervals based on the radiosondes rather than surface based measurements of temperature  $(\theta)$ , humidity (q) and mixed layer depth (h). Whether the substantial differences between the radiosonde and surface based data are due to measurement inaccuracy or a difference in sampling is not known.

The calculation of evolutions of the well mixed  $\theta$  and q variables is shown in Fig. 11. Since the Deardorff model substantially over-estimated  $W_e$  through the analysis period, the model predicted too much warming. This was not only because of the entrainment of too much warm air but also because of the calculated reduction of long wave/cloud-top radiative cooling due to overthinning of the stratus cloud layer by the model. Another factor in the prediction of  $\theta$  and q was the overestimation of the surface fluxes caused by the use of a sea surface temperature higher than the average for the 12 hour period (the sea surface temperature declined by roughly 0.8  $^{\circ}$ C during each 12 hour period).

The tendency of the Deardorff model to overestimate  $W_{\rm e}$  for thin clouds has been discussed by Stage and Businger (1981). They use a slightly different parameterization and find substantially lower values of  $W_{\rm e}$  if  $Z_{\rm c}/h$  is close to one ( $Z_{\rm c}$  is the lifting condensation level). A comparison of  $W_{\rm e}$  estimates of Deardorff (1976) and Stage and Businger (1981) is shown in Table IV.

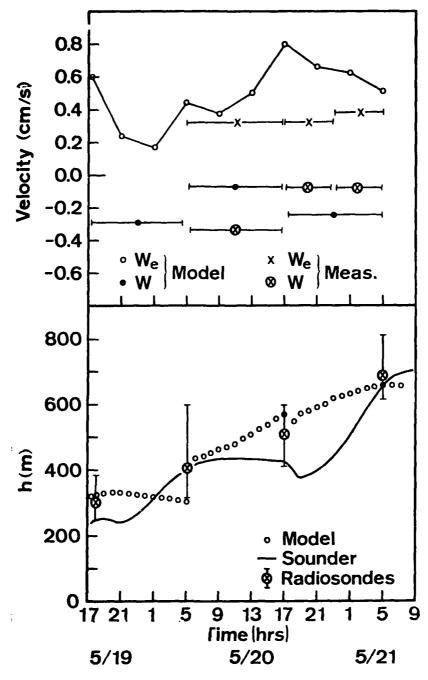


Fig. 10 Entrainment rate (W<sub>e</sub>) and mean vertical motion (W) are on the upper scale and the mixed layer depth (h) is on the lower scale. The symbols are defined in the graphs. The discontinuities in the model results are due to re-initial; zation at 12 hour intervals

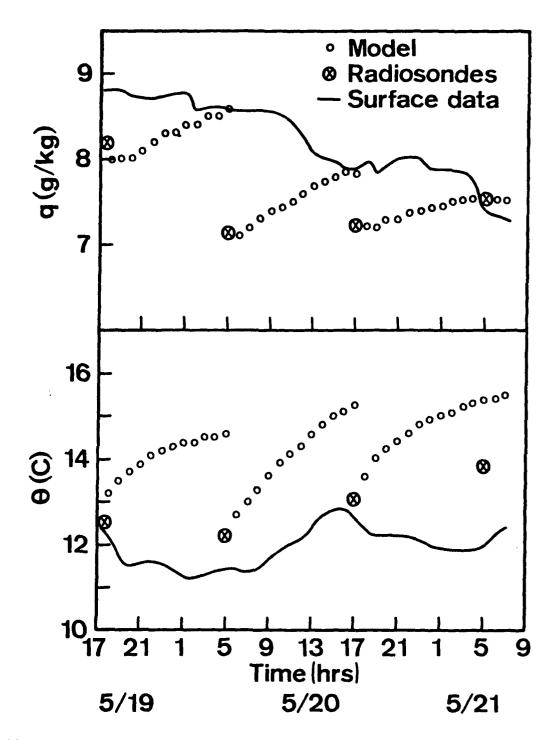


Fig. 11 Temporal evolution of the well mixed meteorological variables  $\theta$  and q. The surface data were measured on the R/V ACANIA

Table IV

A comparison of Deardorff (1976),  $W_e$ , Stage and Businger (1981),  $W_e$ , entrainment rates for the initial conditions at each of the 12-hour analysis periods.

Date	Time	h(m)	$\mathbf{Z}_{\mathbf{C}}(\mathbf{m})$	$z_{c}/h$	W <sub>e</sub> (cm/s)	Wa'(cm/s)
5/19	1700	310	10	z_/h 0.03	Ö. 86	We'(cm/s) 0.86
5/20	0500	<b>38</b> 0	<b>27</b> 0	0.71	0.84	0.42
5/20	1700	<b>52</b> 0	340	0.65	1. <b>4</b> 3	0.29

One final point to consider: the evolution shown in Fig. 9 is calculated at  $S = S_O = 0.8$ . Thus, applications requiring the ambient spectra including the humidity influence (for example, optical extinction) are subject to additional uncertainty due to erroneous model predictions of S (Fig. 12) and therefore g(S). This is particularly critical if S > 0.9.

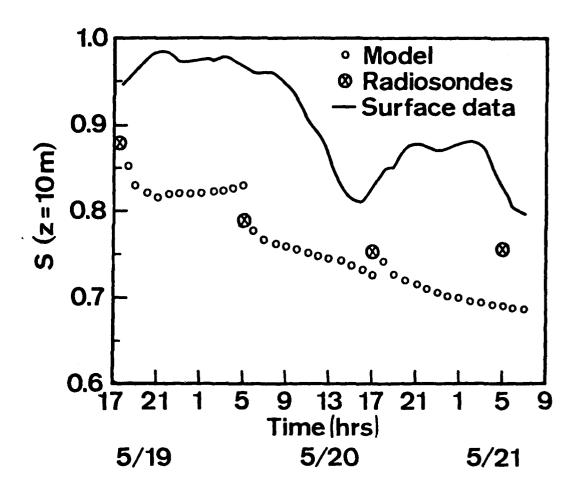


Fig. 12 Similar to Figure 11 but for the ambient saturation ratio S at Z = 10 m

### 8. Conclusions

The application of modern mixed layer physics to the structure and evolutions of atmospheric aerosols in the marine regime represents a philosophical quantum jump over the primarily empirical formulations of the WMK type. This was accomplished by partitioning the aerosol spectrum into continental and locally generated components and by writing the equations in terms of aerosol spectral densities transformed to a predetermined, fixed reference relative humidity. Besides the wind speed and relative humidity, the new formulation requires knowledge or specification of the entrainment rate, the mixed layer height and the sea-surface production rate of droplets. If dynamic effects are ignored, the model can predict an equilibrium aerosol spectrum (Eqn 15) roughly equivalent to that of the WMK model.

The aerosol dynamic model was tested against a marine data set in two ways. In Section 6, Eqn 13a was evaluated under optimum computational circumstances using measurements of the relevant variables wherever possible. In Section 7, a more realistic test was performed where a full mixed layer dynamic model evolution was used to simulate an actual field application of the aerosol model. In both cases the aerosol mixed layer model performed reasonably well but it did not convincingly outperform the much simpler WMK model. The errors in prediction of the aerosol spectrum and the ambient humidity represent about a factor-of-three uncertainty in the aerosol optical extinction coefficient.

Although the mixed layer model was only a modest improvement over WMK model, it holds considerable potential for improvement. The inclusion of advective effects and improvements in the entrainment rate and account surface production flux should be pursued. Another factor to considerable nature of the aerosol production used here. The flux data given

Table III are actually the net surface flux values which represent the droplet production minus losses due to turbulent deposition under average conditions. A modification of Table III to represent the true production and the inclusion of the turbulent deposition velocity (Slinn and Slinn, 1980) explicitly in Eqn 13, is presently under investigation.

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